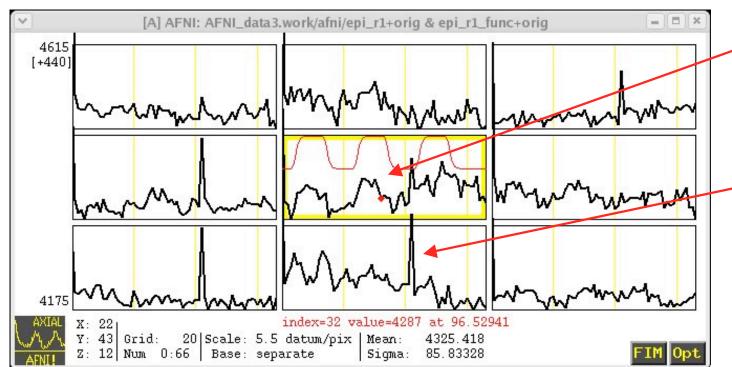
Sample Data Analysis: Simple Regression

- Enough theory (for now: more to come later!)
- To look at the data: type cd AFNI_data3/afni; then afni
- Switch Underlay to dataset epi_r1
 - > Then Axial Image and Graph
 - > FIM-Pick Ideal; then click afni/epi r1 ideal.1D; then Set
 - > Right-click in image, Jump to (ijk), then 22 43 12, then Set



- Data clearly has activity in sync with reference
 - o 30 s blocks
- Data also has a big spike in middle, which is annoying
 - Subject head movement!
 - \circ Also spike at t=0

Preparing Data for Analysis

- Six preparatory steps are common:
 - > Temporal alignment: program <a>3dTshift
 - Image registration (AKA realignment): program 3dvolreg
 - > Image smoothing: program <a>3dmerge
 - Image masking: 3dAutomask or 3dClipLevel
 - Conversion to percentile: programs <u>3dTstat</u> and <u>3dcalc</u>
 - Censoring out time points that are bad: program 3dToutcount (or 3dTqual) and 3dvolreg
- Not all steps are necessary or desirable in any given case
- In this first example, will only do registration, since the data obviously needs this correction

Data Analysis Script

In file epi r1 regress: 3dvolreg (3D image registration) 3dvolreg -base 3 will be covered in detail in a later -verb presentation -prefix epi r1 reg ◆ filename to get estimated motion parameters -1Dfile epi r1 mot.1D epi r1+orig <u>3dDeconvolve</u> = regression code 3dDeconvolve Name of input dataset (from 3dvolreg) -input epi r1 reg+orig -nfirst \ ■ Number of input model time series -num stimts 1 -stim_times 1 epi_r1_times.1D \ → Name of input stimulus class timing file (t's) 'BLOCK (30) ' Name for results in AFNI menus -stim label 1 AllStim \ Indicates to output t-statistic for β weights -tout \ ■ Name of output "bucket" dataset (statistics) -bucket epi r1 func Name of output model fit dataset -fitts epi r1 fitts Name of image file to store X [AKA R] matrix -xjpeg epi r1 Xmat.jpg Name of text file in which to store X matrix -x1D epi r1 Xmat.x1D

Type tcsh epi_r1_regress; then wait for programs to run

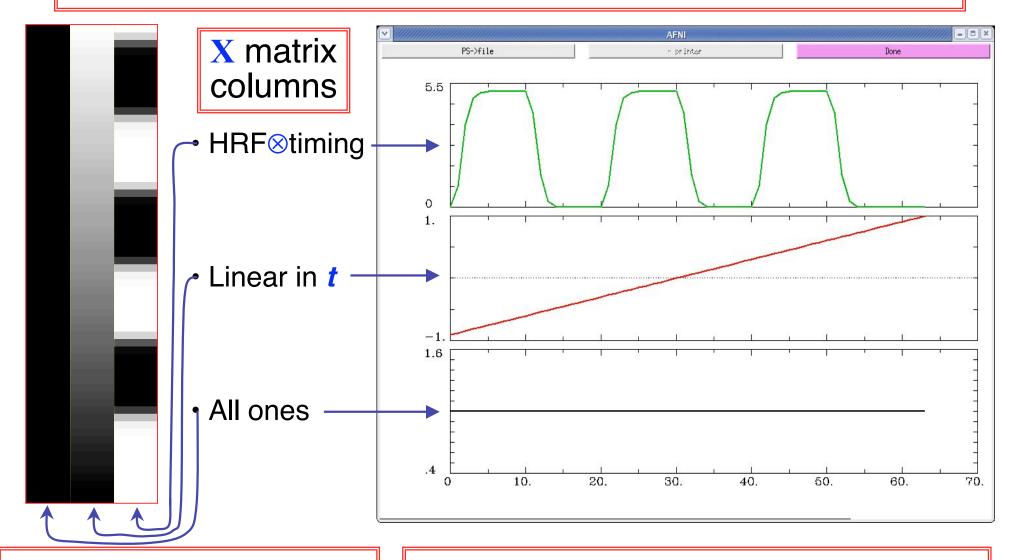
Screen Output of the epi_r1_decon script

 3dvolreg Output ++ 3dvolreg: AFNI version=AFNI 2007 05 29 1644 (Sep 5 2007) [64-bit] ++ Reading input dataset ./epi r1+orig.BRIK ++ Edging: x=3 y=3 z=2++ Initializing alignment base ++ Starting final pass on 67 sub-bricks: 0..1..2..3.. *** ..63..64..65..66.. ++ CPU time for realignment=5.35 s [=0.0799 s/sub-brick] ++ Min : roll=-0.103 pitch=-1.594 yaw=-0.038 dS=-0.354 dL=-0.021 dP=-0.191 ++ Mean: roll=-0.047 pitch=+0.061 yaw=+0.023 dS=+0.006 dL=+0.032 dP=-0.076 ++ Max : roll=+0.065 pitch=+0.290 yaw=+0.055 dS=+0.050 dL=+0.120 dP=+0.113 ++ Max displacement in automask = 2.46 (mm) at sub-brick 42 } Maximum movement estimate ++ Wrote dataset to disk in ./epi r1 reg+orig.BRIK 3dDeconvolve Output ++3dDeconvolve: AFNI version=AFNI 2007 05 29 1644 (Sep 5 2007) [64-bit] ++ Authored by: B. Douglas Ward, et al. ++ loading dataset epi r1 reg+orig *+ WARNING: Input polort=1; Longest run=201.0 s; Recommended minimum polort=2 Consider '-polort 2' ++ -stim times using TR=3 seconds ++ '-stim times 1' using LOCAL times ++ Wrote matrix image to file epi r1 Xmat.jpg ++ Wrote matrix values to file epi_r1_Xmat.x1D Output file indicators ++ Signal+Baseline matrix condition [X] (64x3): 2.59165 ++ VERY GOOD ++ ++ Signal-only matrix condition [X] (64x1): 1 ++ VERY GOOD ++ ++ Baseline-only matrix condition [X] (64x2): 1.08449 ++ VERY GOOD ++ ++ -polort-only matrix condition [X] (64x2): 1.08449 ++ VERY GOOD ++ ++ Matrix inverse average error = 5.62791e-16 ++ VERY GOOD ++ ++ Calculations starting; elapsed time=0.238 ++ voxel loop:0123456789.0123456789.0123456789.0123456789.0123456789.} Progress meter/pacifier ++ Calculations finished; elapsed time=1.417 ++ Wrote bucket dataset into ./epi r1 func+orig.BRIK ++ Wrote 3D+time dataset into ./epi_r1_fitts+orig.BRIK \ Output file indicators ++ #Flops=3.11955e+08 Average Dot Product=4.50251

If a program crashes, we'll need to see this text output (at the very least)!

Stimulus Timing: Input and Visualization

```
epi_r1_times.1D = 9.0 69.0 129.0
= times of start of each BLOCK(30) HRF copy
```



aiv epi_r1_Xmat.jpg

1dplot -sepscl epi_r1_Xmat.x1D

Look at the Activation Map

- Run afni to view what we've got (N.B.: a weak test with only 1 run)
 - Switch Underlay to epi r1 reg (input for 3dDeconvolve)
 - Switch Overlay to epi r1 func (output from 3dDeconvolve)
 - Sagittal Image and Graph viewers
 - > FIM→Ignore→3 to have graph viewer not plot 1st 3 time pts
 - ➤ FIM→Pick Ideal; pick epi_r1_ideal.1D (modeled HRF: output from waver)
- Define Overlay to set up functional coloring
 - \triangleright Olay \rightarrow Allstim#0_Coef (sets coloring to be from model fit β)
 - ➤ Thr→Allstim#0_Tstat (sets threshold to be model fit t-statistic)
 - > See Overlay (otherwise won't see the function!) should be on automatically
 - > Play with threshold slider to get a meaningful activation map (e.g., t=3 is a decent threshold more on thresholds later)
 - > Again, use Jump to (i j k) to jump to index coordinates 22 43 12

More Looking at the Results

- Graph viewer: Opt→Tran 1D→Dataset #N to plot the model fit dataset output by 3dDeconvolve
 - Will open the control panel for the Dataset #N plugin
 - Click first Input line to be 'on'; then choose Dataset epi_r1_reg+orig
 - Also choose Color dk-blue to get a pleasing plot
 - Click 2nd Input on; then choose Dataset epi_r1_fitts+orig
 - Also choose Color limegreen to get a pleasing plot
 - Then click on Set+Close (to close the plugin's control panel)
 - This tool lets you visualize the quality of the data fit
- Can also now overlay function on MP-RAGE anatomical by using Switch Underlay to anat+orig dataset
 - Probably won't want to graph the anat+orig dataset!

Setting the Threshold: Principles

- Bad things (i.e., errors):

 - False negatives non-activations reported where there should be true activations found
 ■ Type II errors
- Usual approach in statistical testing is to control the probability of a type I error (the "p-value")
- In FMRI, we are making many statistical tests: one per voxel (≈20,000+) the result of which is an "activation map":
 - Voxels are colorized if they survive the statistical thresholding process

Setting the Threshold: Bonferroni

- If we set the threshold so there is a 1% chance that any given voxel is declared "active" even if its data is pure noise (FMRI jargon: "uncorrected" p-value is 0.01):
 - And assume each voxel's noise is independent of its neighbors (not really true)
 - With 20,000 voxels to threshold, would expect to get 200 false positives this may be as many as the true activations! Situation: Not so good.
- Bonferroni solution: set threshold (e.g., on t-statistic) so high that uncorrected p-value is 0.05/20000=2.5e-6
 - Then have only a 5% chance that even a single false positive voxel will be reported
 - Objection: will likely lose weak areas of activation

Setting the Threshold: Spatial Clustering

- Cluster-based detection lets us lower the statistical threshold and still control the false positive rate
- Two thresholds:
 - First: a per-voxel threshold that is somewhat low (so by itself leads to a lot of false positives, scattered around)
 - Second: form clusters of spatially contiguous (neighboring) voxels that survive the first threshold, and keep only those clusters above a volume threshold e.g., we don't keep isolated "active" voxels
- Usually: choose volume threshold, then calculate voxel-wise statistic threshold to get the overall "corrected" p-value you want (typically, corrected p=0.05)
 - No easy formulas for this type of dual thresholding, so must use simulation: AFNI program AlphaSim

AlphaSim: Clustering Thresholds

- Simulated for brain mask of 18,465 voxels
- Look for smallest cluster with corrected *p* < 0.05

Uncorrected	Cluster Size	Cluster Size
<i>p</i> -value	/ Corrected p	/ Corrected p
(per voxel)	(uncorrelated)	(correlated 5 mm)
0.0002	2/0.001	3 / 0.004
0.0004	2/0.008	4 / 0.012
0.0007	2 / 0.026	3 / 0.031
0.0010	3 / 0.001	4 / 0.007
0.0020	3 / 0.003	4 / 0.032
0.0030	3 / 0.008	5 / 0.013
0.0040	3 / 0.018	5 / 0.029
0.0050	3 / 0.030	6 / 0.012
0.0060	4 / 0.003	6 / 0.023
0.0070	4 / 0.004	6 / 0.036
0.0080	4 / 0.006	7 / 0.016
0.0090	4 / 0.010	7 / 0.027
0.0100	4 / 0.015	7 / 0.042

Corresponds to sample data

Can make activation maps for display with cluster editing using 3dmerge program – or in AFNI GUI (new: Sep 2006)

End of Important Aside

Multiple Stimulus Classes

- The experiment analyzed here in fact is more complicated
 - There are 9 related communication stimulus types in a 3x3 design of Category by Affect (stimuli are shown to subject as pictures)
 - Telephone, Email & Face-to-face = categories
 - Negative, Positive & Neutral = affects
 - ✓ telephone stimuli: tneg, tpos, tneu
 - ✓ email stimuli: eneg, epos, eneu
 - ✓ face-to-face stimuli: fneg, fpos, fneu
 - > Each stimulus type has 3 presentation blocks of 30 s duration
 - > Scrambled pictures (baseline) are shown between blocks
 - > 9 imaging runs, 64 useful time points in each
 - o Originally, 67 TRs per run, but skip first 3 for MRI signal to reach steady state (i.e., eliminate initial transient spike in data)
 - $_{0}$ So 576 TRs of data, in total (64×9)
 - > Already registered and put together into one dataset: rall_vr+orig

Regression with Multiple Model Files

- Script file rall_decon does the job:
- Run this script by typing tcsh rall_regress (takes a few minutes)

```
3dDeconvolve -input rall vr+orig
                                                            -jobs 2
                                                            -concat '1D: 0 64 128 192 256 320 384 448 512'
   -num stimts 15 -local times
   -stim times 1 '1D: 0 | | 120
                                                'BLOCK(30)'
                                                            \ ← stimulus times
                                         | 120' 'BLOCK(30)'
                                                            -stim times 2 '1D: * | | 120 |
                                                            -stim times 3 '1D: * | 120 | |
                                               'BLOCK(30)'
   -stim times 4 '1D: 60 | | | | 120
                                               'BLOCK (30) '
   -stim times 5 '1D: * | 60 | | 0
                                               'BLOCK(30)'
   -stim times 6 '1D: * | | 0 | |
                                               'BLOCK (30) '
   -stim times 7 '1D: * | 0 | | | 120 |
                                              'BLOCK(30)'
   -stim times 8 '1D: 120 | | |
                                               'BLOCK (30) '
   -stim times 9 '1D: * | | 60 | | | 0 | | 120 | '
                                              'BLOCK(30)'
   -stim label 1 tneg -stim label 2 tpos -stim label 3 tneu
                                                                 → stimulus label
   -stim label 4 eneg -stim label 5 epos -stim label 6 eneu
   -stim label 7 fneg -stim label 8 fpos -stim label 9 fneu
```

continued ...

Regression with Multiple Model Files (continued)

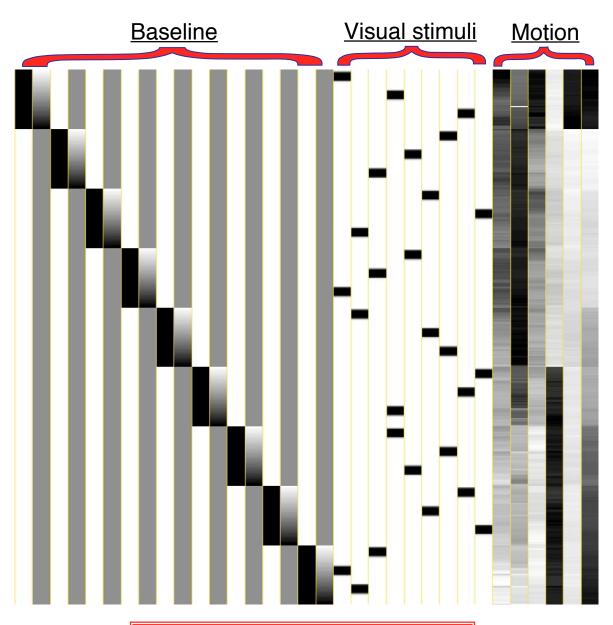
```
-stim file 10 motion.1D'[0]' -stim base 10
                                                  → apply to baseline
-stim file 11 motion.1D'[1]' -stim base 11
-stim file 12 motion.1D'[2]' -stim base 12
-stim file 13 motion.1D'[3]' -stim base 13
-stim file 14 motion.1D'[4]' -stim base 14
-stim file 15 motion.1D'[5]' -stim base 15
-gltsym 'SYM: tpos -epos' -glt label 1 TPvsEP
                                                         → symbolic GLT

→ label the GLT

-gltsym 'SYM: tpos -tneg' -glt label 2 TPvsTNg
-gltsym 'SYM: tpos tneu tneg -epos -eneu -eneg'
        -qlt label 3 TvsE
                                                         → statistic types to output
-fout -tout
-bucket rall func -fitts rall fitts
-xjpeg rall xmat.jpg -x1D rall xmat.x1D
```

- the 9 visual stimulus classes were given using -stim_times
- it is important to include motion parameters as regressors
 - > this helps to exclude stimulus correlated motion artifacts
 - > 6 motion parameters specified as covariates of no interest via
 - -stim_file and -stim_base
 - > 3dvolreg has previously been run, with the -1Dfile option, which gave us file motion. 1D

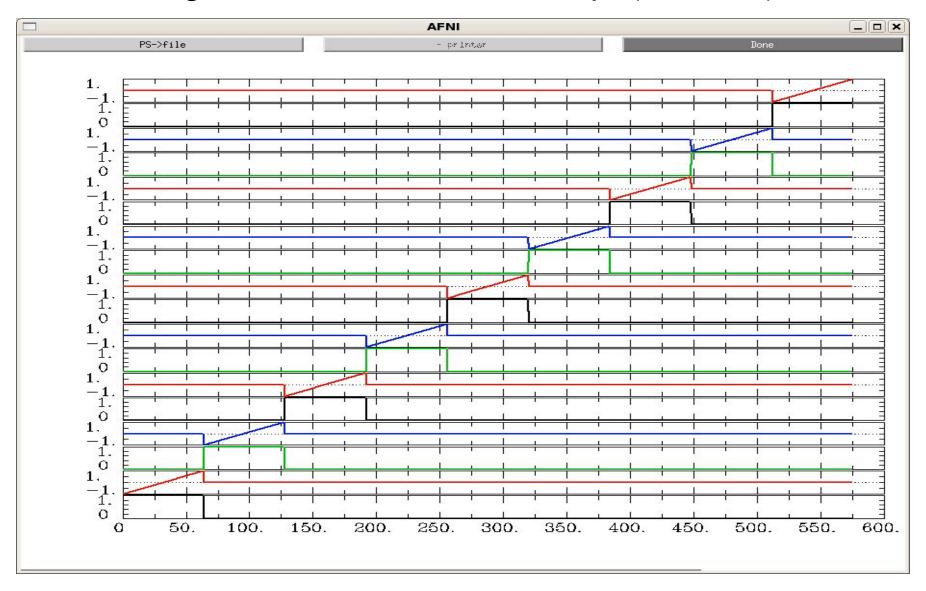
Regressor Matrix for This Script (via -xjpeg)



- 18 baseline regressors
 - > linear baseline
 - > 9 runs times 2 params
- 9 visual stimulus regressors
 - > 3×3 design
- 6 motion regressors
 - > 3 rotations and 3 shifts

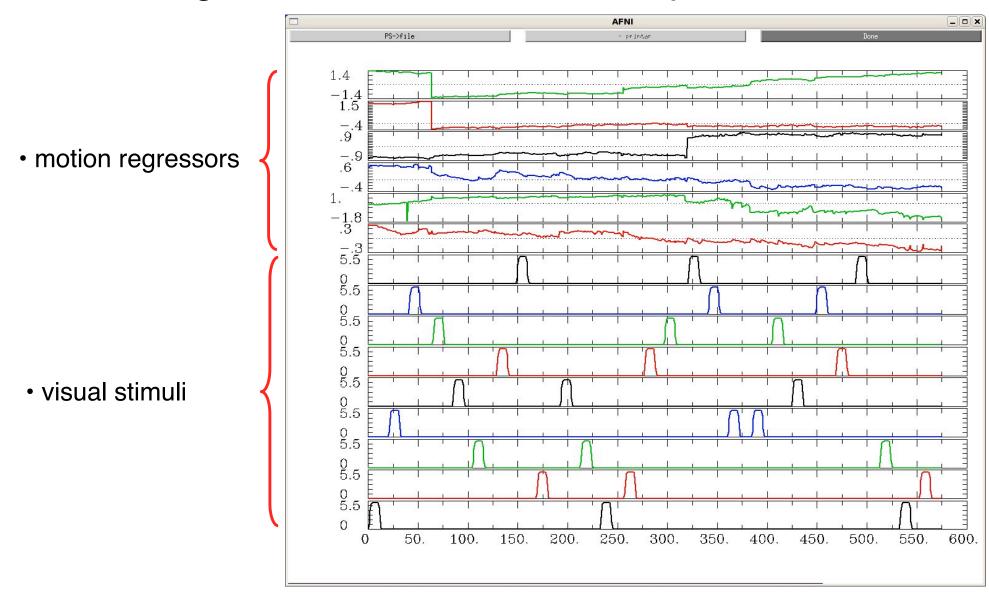
aiv rall_xmat.jpg

Regressor Matrix for This Script (via -x1D)



baseline regressors: via 1dplot -sepscl xmat_rall.x1D'[0..18]'

Regressor Matrix for This Script (via -x1D)



1dplot -sepscl xmat_rall.x1D'[18..\$]'

<u>Options in 3dDeconvolve - 1</u>

```
-concat '1D: 0 64 128 192 256 320 384 448 512'
```

- "File" that indicates where distinct imaging runs start inside the input file
 - Numbers are the time indexes inside the dataset file for start of runs
 - > In this case, a text format .1D file put directly on the command line
 - Could also be a filename, if you want to store that data externally

```
-num stimts 15 -local times
```

- We have 9 visual stimuli (+6 motion), so will need 9 -stim_times below
- Times given in the -stim_times files are local to the start of each run (vs. -global_times meaning times are relative the start of the first run)

```
-stim_times 1

>'1D: 0.0 | | 120.0 | | | | 60.0'

'BLOCK(30,1)'
```

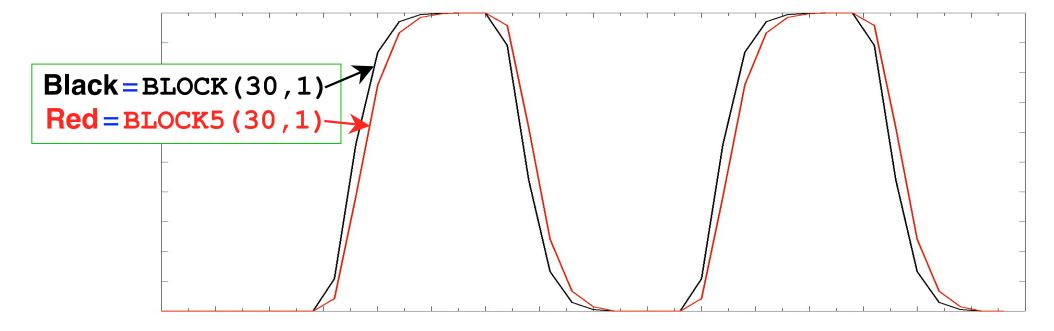
- "File" with 9 lines, each line specifying the start time in seconds for the stimuli within the corresponding imaging run, with the time measured relative to the start of the imaging run itself (local time)
- HRF for each block stimulus is now specified to go to maximum value of 1 (compare to graphs on previous slide)
 - This feature is useful when converting FMRI response magnitude to be in units of percent of the mean

Aside: the 'BLOCK()' HRF Model

• **BLOCK (L)** is convolution of square wave of duration **L** with "gamma variate function" $t^4 e^{-t} / [4^4 e^{-4}]$ (peak value=1 at t=4):

$$h(t) = \int_0^{\min(t,L)} s^4 e^{-s} / [4^4 e^{-4}] ds$$

- "Hidden" option: **BLOCK5** replaces "4" with "5" in the above
 - Slightly more delayed rise and fall times
- BLOCK (L, 1) makes peak amplitude of block response = 1



Options in 3dDeconvolve - 2

```
-gltsym 'SYM: tpos -epos' -glt label 1 TPvsEP
```

- GLTs are General Linear Tests
- 3dDeconvolve provides test statistics for each regressor separately, but if you want to test combinations or contrasts of the β weights in each voxel, you need the -gltsym option
- Example above tests the difference between the β weights for the Positive Telephone and the Positive Email responses
 - Starting with SYM: means symbolic input is on command line
 Otherwise inputs will be read from a file
 - > Symbolic names for each regressor taken from -stim_label options
 - Stimulus label can be preceded by + or to indicate sign to use in combination of β weights
 - Leave space after each label!
- Goal is to test a linear combination of the β weights
 - Tests if $\beta_{tpos} = \beta_{epos}$
 - e.g., does tpos get different response from epos ?
- Quiz: what would 'SYM: tpos epos' test?

 $0 = soq_b + soq_b$ if test bluow th

Options in 3dDeconvolve - 3

```
-gltsym 'SYM: tpos tneu tneg -epos -eneu -eneg'
-glt_label 3 TvsE
```

- Goal is to test if $(\beta_{tpos} + \beta_{tneu} + \beta_{tneg}) (\beta_{epos} + \beta_{eneu} + \beta_{eneg}) = 0$
 - Test average BOLD signal change among the 3 affects in the telephone tasks versus the email tasks

```
-gltsym 'SYM: tpos -epos | tneu -eneu | tneg -eneg'
-glt_label 3 TvsE_F
```

- Goal is to test if $\beta_{tpos} = \beta_{epos}$, $\beta_{tneu} = \beta_{eneu}$, and $\beta_{tneg} = \beta_{eneg}$ are all true
 - BOLD signal change of any affect in the telephone tasks versus the email tasks
 - This is a different test than the previous one!
- -glt_label 3 TvsE option is used to attach a meaningful label to the resulting statistics sub-bricks
 - Output includes the ordered summation of the β weights and the associated statistical parameters (t- and/or F-statistics)
 - t- or F-statistics?

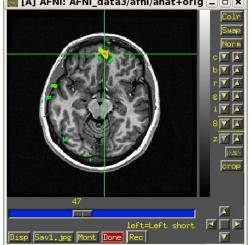
Options in 3dDeconvolve - 4

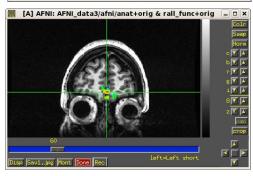
-fout -tout

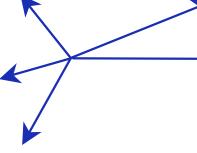
- = output both F- and t-statistics for each stimulus class (-fout) and stimulus coefficient (-tout) — but not for the baseline coefficients (if you want baseline statistics: -bout)
- The full model statistic is an *F*-statistic that shows how well the sum of all 9 input model time series fits voxel time series data
 - ➤ Compared to how well *just* the baseline model time series fit the data times (in this example, have 24 baseline regressor columns in the matrix 18 for the linear baseline, plus 6 for motion regressors)
 - \rightarrow F = [SSE(r) SSE(f)]/df(n) ÷ [SSE(f)/df(d)]
- The individual stimulus classes also will get individual F- and/or t-statistics indicating the significance of their individual incremental contributions to the data time series fit
 - > e.g., F_{tpos} (#6, equivalent to t (#5)) tells if the full model explains more of the data variability than the model with tpos omitted and all other model components included

Results of rall_regress Script

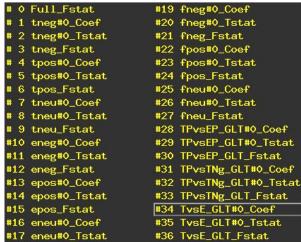








#18 eneu_Fstat



 Images showing results from third GLT contrast: TvsE

- Menu showing labels
 from 3dDeconvolve run
- Play with these results yourself!

Statistics from 3dDeconvolve

- An F-statistic measures significance of how much a model component (stimulus class) reduced the variance (sum of squares) of data time series residual
 - After all the other model components were given their chance to reduce the variance
 - > Residuals = data model fit = errors = -errts
 - > A *t*-statistic sub-brick measures impact of one coefficient (of course, **BLOCK** has only one coefficient)
- Full F measures how much the all regressors of interest combined reduced the variance over just the baseline regressors (sub-brick #0)
- Individual partial-model Fs measures how much each individual signal regressor reduced data variance over the full model with that regressor excluded (e.g., sub-bricks #3, #6, #9)
- The Coef sub-bricks are the β weights (e.g., #1, #4, #7, #10) for the individual regressors
- Also present: GLT coefficients and statistics

0 Full_Fstat #19 fneg#0_Coef # 1 tneg#0_Coef #20 fneg#0_Tstat # 2 tneg#0_Tstat #21 fneg_Fstat # 3 tneg_Fstat #22 fpos#0_Coef # 4 tpos#0_Coef #23 fpos#0_Tstat # 5 tpos#0_Tstat #24 fpos_Fstat # 6 tpos_Fstat #25 fneu#0_Coef # 7 tneu#O_Coef #26 fneu#0_Tstat # 8 tneu#0_Tstat #27 fneu_Fstat # 9 tneu Fstat #28 TPvsEP_GLT#O_Coef #10 eneg#0_Coef #29 TPvsEP_GLT#0_Tstat #11 eneg#0_Tstat #30 TPvsEP_GLT_Fstat #12 eneg_Fstat #31 TPvsTNg_GLT#0_Coef #13 epos#0_Coef #32 TPvsTNg_GLT#0_Tstat #33 TPvsTNg_GLT_Fstat #14 epos#0_Tstat #34 TvsE_GLT#0_Coef #15 epos_Fstat #16 eneu#0_Coef #35 TvsE_GLT#0_Tstat #17 eneu#0_Tstat #36 TvsE_GLT_Fstat #18 eneu_Fstat

Group Analysis: will be carried out on β or **GLT** coefs from single-subject analyses